

09/877.748

09/877, 748

p.1

Individual Applicant

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City : Rochester
State : New York
Country : USA
PostalCode : 14620-____
PhoneNumber : ____-____-____
FaxNumber : ____-____-____
EmailAddress :

<110> LastName : Sutherland
<110> FirstName : John
<110> MiddleInitial : W
<110> Suffix :

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EmailAddress :

<110> LastName : Sutherland
<110> FirstName : John
<110> MiddleInitial : W
<110> Suffix :

Application Project

<120> Title : Detecting Nucleic Acid Detection Sequences
<130> AppFileReference : CDS-232
<140> CurrentAppNumber :
<141> CurrentFilingDate : ____-____-____

Sequence

<213> OrganismName : Synthetic construct
<400> PreSequenceString :
cacagacatc ataacaaaaa atttccacca aacccccccct ccccgcttc tggccacagc
60acttaaacac atctctgcc aacccccaaaa acaaagaacc ctaacac
<212> Type : DNA
<211> Length : 107
SequenceName : First Sequence
SequenceDescription :

Custom Codon

Sequence Name : First Sequence

Does Not Comply
Corrected Diskette Needed

Non valid sequence
listing format. See
attached example of
valid sequence listing
and explanation of
features.

APPENDIX 3

SPECIMEN SEQUENCE LISTING

<110> Smith, John; Smithgene Inc.

<120> Example of a Sequence Listing

<130> 01-00001

<140> PCT/EP98/00001

<141> 1998-12-31

<150> US 08/999,999

<151> 1997-10-15

<160> 4

<170> PatentIn version 2.0

<210> 1

<211> 389

<212> DNA

<213> Paramecium sp.

<220>

<221> CDS

<222> (279)...(389)

<300>

<301> Doe, Richard

<302> Isolation and Characterization of a Gene Encoding a
Protease from Paramecium sp.

<303> Journal of Genes

<304> 1

<305> 4

<306> 1-7

<307> 1988-06-31

<308> 123456

<309> 1988-06-31

<400> 1

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tgatgtggca	attgctggca	gtgccacagg	cttttcagcc	aggcttaggg	tgggttcgcg	180
cgcggcgcg	cggccctct	cgcgctcctc	tcgcgcctct	ctctcgctct	cctctcgctc	240

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ggacctgatt	aggtgagcag	gaggaggggg	cagtttagc	atg Met 1	ggt Val	tca Ser	atg Met	ttc Phe 5	agc Ser	296						
ttg Leu	tct Ser	ttc Phe	aaa Lys 10	tgg Trp	cct Pro	gga Gly	ttt Phe	tgt Cys 15	ttg Leu	ttt Phe	ggt Val	tgt Cys	ttg Leu 20	ttc Phe	caa Gln	344
tgt Cys	ccc Pro	aaa Lys 25	gtc Val	ctc Leu	ccc Pro	tgt Cys	cac His 30	tca Ser	tca Ser	ctg Leu	cag Gln	ccg Pro 35	aat Asn	ctt Leu	389	
<210>	2															
<211>	37															
<212>	PRT															
<213>	Paramecium sp.															
<400>	2															
Met 1	Val	Ser	Met	Phe 5	Ser	Leu	Ser	Phe	Lys 10	Trp	Pro	Gly	Phe	Cys 15	Leu	
Phe	Val	Cys	Leu 20	Phe	Gln	Cys	Pro	Lys 25	Val	Leu	Pro	Cys	His 30	Ser	Ser	
Leu	Gln	Pro 35	Asn	Leu												
<210>	3															
<211>	11															
<212>	PRT															
<213>	Artificial Sequence															
<220>																
<223>	Designed peptide based on size and polarity to act as a linker between the alpha and beta chains of Protein XYZ.															
<400>	3															
Met 1	Val	Asn	Leu	Glu 5	Pro	Met	His	Thr	Glu 10	Ile						
<210>	4															
<400>	4															
000																

[Annex VIII follows]

Mandatory (M) or optional (O).		Comments and format	
<110>	Applicant	Preferably max. of 10 names; one name preferable format: Surname, Other Names and/or Initials.	M. <i>Note</i>
<120>	Title of Invention		M.
<130>	File Reference	Personal file reference	M when filed prior to assignment of appl. number.
<140>	Current Application Number.	Specify as: US 07/999,999 or PCT/US96/99999	M, if available.
<141>	Current Filing Date	Specify as: yyyy-mm-dd	M, if available.
<150>	Prior Application Number.	Specify as: US 07/999,999 or PCT/US96/99999	M, if applicable include priority documents under 35 USC 119 and 120.
<151>	Prior Application Filing Date.	Specify as: yyyy-mm-dd	M, if applicable.
<160>	Number of SEQ ID NOs	Count includes total number of SEQ ID NOs	M.
<170>	Software	Name of software used to create the Sequence Listing.	O.
<210>	SEQ ID NO:1	Response shall be an integer representing the SEQ ID NO shown.	M.
<211>	Length	Respond with an integer expressing the number of bases or amino acid residues.	M.

Numeric Identifier	Definition	Comments and format	Mandatory (M) or optional (O).
<212>	Type	Whether presented sequence molecule is DNA, RNA, or PRT (protein). If a nucleotide sequence contains both DNA and RNA fragments, the type shall be "DNA." In addition, the combined DNA/RNA molecule shall be further described in the <220> to <223> feature section.	M.
<213>	Organism	Scientific name, i.e. Genus/ species, Unknown or Artificial Sequence. In addition, the "Unknown" or "Artificial Sequence" organisms shall be further described in the <220> to <223> feature section.	M
<220>	Feature	Leave blank after <220>. <221-223> provide for a description of points of biological significance in the sequence.	M, under the following conditions: if "n," "Xaa," or a modified or unusual L-amino acid or modified base was used in a sequence; if ORGANISM is "Artificial Sequence" or "Unknown"; if molecule is combined DNA/RNA
<221>	Name/Key	Provide appropriate identifier for feature, preferably from WIPO Standard ST.25 (1998), Appendix 2, Tables 5 and 6.	M, under the following conditions: if "n," "Xaa," or a modified or unusual L-amino acid or modified base was used in a sequence.
<222>	Location	Specify location within sequence; where appropriate state number of first and last bases/ amino acids in feature.	M, under the following conditions: if "n," "Xaa," or a modified or unusual L-amino acid or modified base was used in a sequence.
<223>	Other Information	Other relevant information; four lines maximum	M, under the following conditions: if "n," "Xaa," or a modified or unusual L-amino acid or modified base was used in a sequence; if ORGANISM is "Artificial Sequence" or "Unknown"; if molecule is combined DNA/RNA.
<300>	Publication Information	Leave blank after <300>	O.
<301>	Authors	Preferably max of ten named authors of publication; specify one name per line; preferable format: Surname, Other Names and/or Initials.	O.
<302>	Title		O.
<303>	Journal		O.
<304>	Volume		O.
<305>	Issue		O.
<306>	Pages		O.
<307>	Date	Journal date on which data published; specify as yyyy-mm-dd, MMM-yyyy or Season-yyyy.	O.
<308>	Database Accession Number.	Accession number assigned by database including database name.	O.
<309>	Database Entry Date	Date of entry in database; specify as yyyy-mm-dd or MMM-yyyy.	O.
<310>	Patent Document Number.	Document number; for patent-type citations only. Specify as, for example, US 07/999,999.	O.
<311>	Patent Filing Date	Document filing date, for patent-type citations only; specify as yyyy-mm-dd.	O.
<312>	Publication Date	Document publication date, for patent-type citations only; specify as yyyy-mm-dd.	O.
<313>	Relevant Residues	FROM (position) TO (position)	O.
<400>	Sequence	SEQ ID NO should follow the numeric identifier and should appear on the line preceding the actual sequence.	M.

```

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accaaaaatga acgaaaatct gttcgcttca ttcatgccc ccacaatcct aggcctaccc 240
gccgcagtag tgatcattct atttccccct ctattgatcc ccacctcaa atatctcatc 300
aacaaccgac taatcaccac ccaacaatga 330

```

```

<210> 6
<211> 27
<212> DNA
<213> Synthetic construct

```

```

<400> 6
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```

```

<210> 7
<211> 21
<212> DNA
<213> Synthetic construct

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<400> 7
tcattgttgg gtggtgatta g 21

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```

<210> 8
<211> 35
<212> DNA
<213> Synthetic construct

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<400> 8
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<210> 9
<211> 28
<212> DNA
<213> Synthetic construct

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<400> 9
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<210> 10
<211> 20
<212> DNA
<213> Synthetic construct

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<400> 10
gcccacaact aatactaccg 20

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<210> 11
<211> 26
<212> DNA
<213> Synthetic construct

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<400> 11
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<210> 12
 <211> 46
 <212> DNA
 <213> Synthetic construct

<400> 12
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<210> 13
 <211> 181
 <212> DNA
 <213> Human

<400> 13
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 ttctgacatt actgccagcc accatgaata ttgtacggtta ccataaatac ttgaccacct 120
 gtagtacata aaaacccaat ccacatcaaa accccctccc catgcttaca agcaagtaca 180
 g 181

<210> 14
 <211> 22
 <212> DNA
 <213> Synthetic construct

<400> 14
 ggggaagcag atttgggtac ca 22

<210> 15
 <211> 20
 <212> DNA
 <213> Synthetic construct

<400> 15
 ctgtacttgc ttgtaagcat 20

<210> 16
 <211> 36
 <212> DNA
 <213> Synthetic construct

<400> 16
 gcgtcggact caccatcaa caaccgctat cgacgc 36

<210> 17
 <211> 198
 <212> DNA
 <213> Human

<400> 17

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gccgcagtac tgatcattct atttcccct ctattgatcc ccacctccaa atatctcatc      60
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taaatacttg accacctgta gtacataaaa acccaatcca catcaaaacc ccctcccat      180
gcttacaagc aagtacag                                     198

```

```

<210> 18
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```

<400> 18
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```

```

<210> 19
<211> 33
<212> DNA
<213> Synthetic construct

```

```

<400> 19
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```

```

<210> 20
<211> 47
<212> DNA
<213> Synthetic construct

```

```

<400> 20
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```

```

<210> 21
<211> 30
<212> DNA
<213> Synthetic construct

```

```

<400> 21
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```

```

<210> 22
<211> 30
<212> DNA
<213> Synthetic construct

```

```

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gaacgaaaat ctgttcgctt cattcattgc                                     30

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```

<210> 23
<211> 39
<212> DNA
<213> Synthetic construct

```

```

<400> 23
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```

<210> 24
<211> 22
<212> DNA
<213> Synthetic construct

<400> 24
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<210> 25
<211> 53
<212> DNA
<213> Synthetic construct

<400> 25
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<210> 26
<211> 43
<212> DNA
<213> Synthetic construct

<400> 26
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